

Institut für Lebensmittelsicherheit und -hygiene
der Vetsuisse-Fakultät, Universität Zürich

Direktor: Prof. Dr. Roger Stephan

Arbeit unter wissenschaftlicher Betreuung von
PD Dr. Herbert Hächler

**Shiga toxin-producing Escherichia coli O157
associated with human infections in Switzerland:
2000-2009**

Inaugural-Dissertation

zur Erlangung der Doktorwürde der
Vetsuisse-Fakultät Universität Zürich

vorgelegt von

Ursula Elisabeth Käppeli

Tierärztin
von Wohlen, Aargau

genehmigt auf Antrag von

Prof. Dr. Roger Stephan, Referent

2010

Meinen lieben Eltern

INHALTSVERZEICHNIS

1. Zusammenfassung	2
1.1 Deutsch	2
1.2 Englisch	3
2. Publikation	
3. Danksagung	12
4. Curriculum Vitae	

1. Zusammenfassung

1.1 Deutsch

Vetsuisse-Fakultät Universität Zürich (2010)

Ursula Käppeli

Institut für Lebensmittelsicherheit und -hygiene, ils@fsafety.uzh.ch

Shiga toxin bildende *Escherichia coli* O157 im Zusammenhang mit humanen Infektionen in der Schweiz: 2000-2009

Zusammenfassung:

Shiga toxin bildende *Escherichia coli* (STEC), wichtige durch Lebensmittel übertragbare Erreger, können milde bis schwere blutige Durchfälle (BD), manchmal gefolgt von lebensbedrohlichen Komplikationen wie dem Hämolytisch-urämisches Syndrom (HUS), verursachen. Insgesamt wurden 44 STEC O157 Stämme, die von verschiedenen Patienten in den Jahren 2000 bis 2009 in der Schweiz isoliert wurden, weiter charakterisiert und die dazugehörigen anamnestischen Daten aufgearbeitet. 15.9% der Patienten hatten nicht blutigen Durchfall, 61.4% der Patienten zeigten BD und 29.5% entwickelten ein HUS. Alle Stämme gehörten zum MLST Typ 11, waren *stx2*, *eae* und *ehxA* positiv und nur zwei Stämme zeigten Antibiotikaresistenzen. Unter den 44 Stämmen wurden 9 Phagen-Typen (PT) nachgewiesen, die häufigsten waren; PT 32 (43.2%) und PT 8 (18.2%). Die Genotypisierung (Pulsfeld Gelelektrophorese) ergab 39 verschiedene Muster. Die hohe genetische Vielfalt innerhalb der Stämme führt zu der Schlussfolgerung, dass STEC O157 Infektionen in der Schweiz häufig als Einzelfälle und nicht als grössere Ausbrüche aufgetreten sind.

Stichworte: Epidemiologie, Infektion beim Menschen, O157, STEC

1.2 Englisch

Vetsuisse Faculty University of Zurich (2010)

Ursula Käppeli

Institute for Food Safety and Hygiene, ils@fsafety.uzh.ch

Shiga toxin-producing *Escherichia coli* O157 associated with human infections in Switzerland: 2000-2009

Summary:

Shiga toxin-producing *Escherichia coli* (STEC), an important foodborne pathogen, can cause mild up to severe bloody diarrhoea (BD), sometimes followed by life-threatening complications such as haemolytic uraemic syndrome (HUS). A total of 44 STEC O157 strains isolated from different patients from 2000 through 2009 in Switzerland were further characterized and linked to anamnestic data. Non-bloody diarrhoea was experienced by 15.9%, BD by 61.4% of the patients, and 29.5% developed HUS. All strains belonged to MLST type 11, were positive for *stx2*, *eae* and *ehxA* and only two strains showed antibiotic resistances. Among the 44 strains, 9 phage types (PTs) were detected, the most frequent ones being PT 32 (43.2%) and PT 8 (18.2%). By pulsed-field gel electrophoresis (PFGE), 39 different patterns were found. This high genetic diversity within the strains leads to the conclusion that STEC O157 infections in Switzerland most often occur as single and sporadic cases.

Key words: epidemiology, human infection, O157, STEC

Shiga toxin-producing *Escherichia coli* O157 associated with human infections in Switzerland, 2000–2009

U. KÄPPELI¹, H. HÄCHLER¹, N. GIEZENDANNER¹, T. CHEASTY²
AND R. STEPHAN^{1*}

¹ Institute for Food Safety and Hygiene, Vetsuisse Faculty, University of Zurich, Zurich, Switzerland

² Laboratory of Gastrointestinal Pathogens, GEZI (HPA Centre for Infections), London, UK

(Accepted 25 August 2010)

SUMMARY

Shiga toxin-producing *Escherichia coli* (STEC), an important foodborne pathogen, can cause mild to severe bloody diarrhoea (BD), sometimes followed by life-threatening complications such as haemolytic uraemic syndrome (HUS). A total of 44 O157 strains isolated from different patients from 2000 through 2009 in Switzerland were further characterized and linked to medical history data. Non-bloody diarrhoea was experienced by 15·9%, BD by 61·4% of the patients, and 29·5% developed HUS. All strains belonged to MLST type 11, were positive for *stx2* variants (*stx2* and/or *stx2c*), *eae* and *ehxA*, and only two strains showed antibiotic resistance. Of the 44 strains, nine phage types (PTs) were detected the most frequent being PT32 (43·2%) and PT8 (18·2%). By PFGE, 39 different patterns were found. This high genetic diversity within the strains leads to the conclusion that STEC O157 infections in Switzerland most often occur as sporadic cases.

Key words: Characteristics, epidemiology, human infection, STEC O157.

INTRODUCTION

Escherichia coli, a commensal of the intestinal tract of humans, as well as many animals, is a Gram-negative, facultative anaerobic rod. The virulent, disease-causing *E. coli* can be divided into two major groups: gastrointestinal and extragastrointestinal pathogens. The gastrointestinal category includes enterotoxigenic *E. coli* (ETEC), enteropathogenic *E. coli* (EPEC), enteroinvasive *E. coli* (EIEC), enteroaggregative *E. coli* (EAEC) and Shiga toxin-producing *E. coli* (STEC), which are significant causes of foodborne diseases. STEC can lead to severe illness, like haemolytic

uraemic syndrome (HUS), which presents with haemolytic anaemia, thrombocytopenia and renal insufficiency, most often after an onset of diarrhoea (bloody or non-bloody). In many different O:H serotypes, there is a group of the so-called top five serogroups (O26, O103, O111, O145, O157) which are mainly related to severe illness in humans. Although the importance of non-O157 serovars is being increasingly recognized, so far O157 has caused most outbreaks and cases of severe disease throughout the world [1–3].

A stepwise evolutionary model postulates that the O157 clonal group splits into one lineage, leading to the common non-sorbitol-fermenting (nSF) O157:H7 clonal complex, and a second branch of sorbitol-fermenting (SF) O157:H[−] (non-motile) strains [4]. Recently, SF STEC O157:H[−] strains have been increasingly isolated from clinical cases [5, 6],

* Author for correspondence: Professor R. Stephan, Institute for Food Safety and Hygiene, Vetsuisse Faculty University of Zurich, Winterthurerstr. 272, CH-8057 Zurich, Switzerland.
(Email: stephanr@fsafety.uzh.ch)

emphasizing the importance of analysing patients' stool samples for both SF and nSF *stx*-producing *E. coli* O157.

E. coli O157:H7/H⁻ exist in humans and in the environment as *stx*-positive and *stx*-negative variants [6, 7]. Friedrich *et al.* [6] compared the prevalence of *stx*-negative strains in SF and nSF *E. coli* O157 isolated from stools of patients with sporadic cases of diarrhoea or HUS. They found that the proportion of *stx*-negative strains was significantly higher in SF O157:H⁻ isolates (12.7%) than in nSF O157:H7/H⁻ isolates (0.8%). These authors also identified *stx*-negative SF *E. coli* O157 strains in patients' stool samples during three outbreaks of HUS and one outbreak of diarrhoea and documented loss of *stx* genes during outbreaks of human infection [6].

Since few specific data of clinical human O157 STEC strains are available in Switzerland, the aim of this study was (i) to further characterize all O157 STEC strains sent to the National Centre for Enteropathogenic Bacteria (NENT) from 2000 to 2009, and (ii) to compare the results with data from other countries.

METHODS

Strains

A total of 144 STEC strains collected and stored (−80 °C) by NENT were used for further characterization. The strains were isolated between 2000 and 2009 from human faecal samples, collected by hospitals or family doctors. All strains were confirmed to be STEC by detection of *stx* genes by polymerase chain reaction (PCR) with primers VT1 and VT2, based on sequence targeting of a conserved region between *stx1* and *stx2* genes and PCR conditions described previously [8].

Serotyping

O157 strains were distinguished from the collection of the 144 STEC strains using PCR with primers based on sequences of the *rfbE* (*per*) gene [9]. STEC O157:H7 strain EDL 933 was included as a control. The H antigen of the O157 strains was determined by PCR with primers FLICH7-F and FLICH7-R, and PCR conditions as described previously [10]. Moreover, all O157 strains were serotyped with an O157:H7 latex agglutination test (Wellcolex *E. coli* O157:H7; Remel, USA).

Further strain characterization

Fermentation of sorbitol was detected on sorbitol MacConkey agar (SMAC) (Oxoid Ltd, UK). Strains were further tested by PCR for *stx1* and *stx2* [11], *eae* and *eae* γ 1 encoding intimin [12], and *ehxA* encoding EHEC haemolysin [13]. Further characterization of the Shiga toxin type 2 variant B-subunit was done by PCR–RFLP [14].

Phage-typing

Bacterial phage-typing was performed at the Laboratory of Gastrointestinal Pathogens, GEZI (HPA Centre for Infections, London, UK) by the methods described by Khakhria *et al.* [15].

Multi-locus sequence typing (MLST)

Internal amplicons of seven housekeeping genes (*adh*, *fumC*, *gyrB*, *icdF*, *mdh*, *purA*, *recA*) were sequenced [16] and alleles as well as sequence types (ST) were assigned in accordance with the *E. coli* MLST website (<http://mlst.ucc.ie/mlst/dbs/Ecoli>).

Genotyping

Pulsed-field gel electrophoresis (PFGE) was performed according to the CDC PulseNet protocol (<http://www.cdc.gov/pulsenet/protocols.htm>) with minor modifications. Briefly, strains were grown on blood agar at 37 °C overnight. Colonies from blood agar were resuspended in cell suspension buffer (OD₆₀₀ = 1). The bacterial cell suspension was mixed with 400 μ l of 1.4% Bio-Rad agarose (Bio-Rad, Germany) and cells were lysed by proteinase K treatment overnight. After lysis the plugs were washed twice for 15 min in ultrapure water and four times for an hour in Tris-EDTA (TE) buffer. After washing with TE buffer, DNA agarose plugs were incubated overnight in the presence of *Xba*I (Roche, Germany) according to the manufacturer's instructions. Restricted DNA in plug slices was separated in 1% SeaKem Gold (BioConcept, Switzerland) agarose gel at 6 V/cm in 0.5× Tris-borate-EDTA buffer cooled to 14 °C in a CHEF-DR III system (Bio-Rad). The pulse times were ramped from 5 s to 50 s for 20 h at an angle of 120°. Gels were stained with ethidium bromide and visualized under UV light transillumination with Gel Doc (Bio-Rad) and analysed with BioNumerics software (Applied Maths, Belgium).

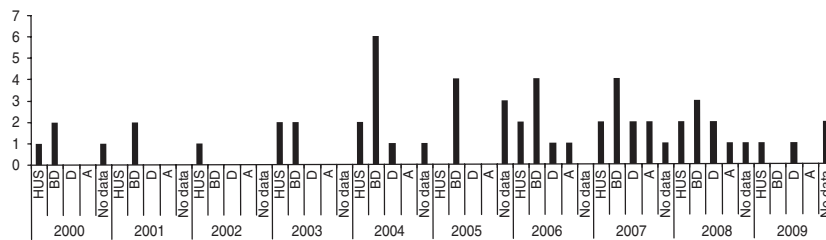


Fig. 1. Medical history data of 44 STEC O157 strains isolated from human patients from 2000 to 2009 in Switzerland. HUS, Haemolytic uraemic syndrome; BD, bloody diarrhoea; D, non-bloody diarrhoea; A, anaemia.

As a reference *Salmonella* Braenderup strain H9812 (ATCC BAA 664) was used.

Antimicrobial susceptibility testing

The strains were tested for antimicrobial resistance by the disk diffusion method according to the protocols of the Clinical Laboratory Standards Institute (CLSI, 2008). The panel of antibiotics (disks: Becton, Dickinson, USA) consisted of ampicillin (AM), amoxicillin/clavulanic acid (AMC), ceftazidime (CAZ), cefalothine (CF), ciprofloxacin (CIP), cefpodoxime (CPD), cefotaxime (CTX), cefuroxime (CXM), cefepime (FEP), ceftazidime (FOX), gentamicin (GM) and tetracycline (Te). *E. coli* ATCC 25922 was used as a quality control strain. The strains were classified as resistant or susceptible to each antibiotic agent. Strains giving 'intermediate' values were considered susceptible.

RESULTS

Medical history data

Of the 144 STEC strains collected during 2000–2009, 44 strains from 44 different patients were identified as O157. Medical history data for these patients are summarized in Figure 1. Thirteen patients (29.5%) developed HUS, of which six (46.2%) were male, and seven (53.8%) female, with an average age of 3.5 years (range 1–15 years). Twenty-seven (61.4%) patients presented with bloody diarrhoea, seven (15.9%) with non-bloody diarrhoea, and four (9.1%) were anaemic. For nine patients no medical history data was available. Thirty-three (76.7%) patients were aged ≤ 10 years, three (7%) were aged ≥ 60 years and seven (16.3%) patients were aged between 10 and 60 years. For one patient (female, presenting with HUS, diarrhoea and anaemia) no data was available.

Further strain characterization

Latex agglutination of all 44 strains was positive for O157, whereas 11 (25%) strains were negative for H7 agglutination (O157:NM). Results are listed in Table 1. Nevertheless, these strains were positive for the *fliC* gene by PCR.

Four (9.1%) strains fermented sorbitol (SF) on SMAC, 40 (90.9%) were nSF. All four SF strains were O157:H⁻, 33 (82.5%) of the nSF strains were O157:H7, and seven (17.5%) strains were O157:H⁻. All 44 *E. coli* O157 strains tested positive for *eae* and *ehxA* genes. Among the strains, 52.3%, 63.6%, 15.9%, and 20.5% harboured *stx1*, *stx2*, *stx2c*, and the combination of *stx2* and *stx2c*, respectively (Table 1).

Phage-typing

Nine different phage types were found (PT2, 4, 8, 14, 23, 32, 49, 50, 71), and six (13.6%) strains could not be further characterized by phage-typing (RDNC, Reacts but Does Not Conform to a published typing pattern). PT2, 4, 14, 50 and 71 appeared only once. PT8 was found eight times (18.2%), PT23 twice (4.5%), PT32 nineteen times (43.2%) and PT49 resulted in four (9.1%) of the strains (Table 1).

MLST

All 44 strains belonged to ML ST 11, similar to the vast majority of *eae*-positive human STEC O157 isolates.

PFGE typing

PFGE patterns were very heterogeneous except for eight strains (18.2%). The remaining 36 patterns showed similarity coefficients between 59% and 94% (Dice similarity index and UPGMA method). The dendrogram is displayed in Figure 2.

Table 1. Serotypes and virulence factors of STEC O157 strains isolated from human patients from 2000 to 2009 in Switzerland

Strain no.	Serotype	Sorbitol	<i>stx1</i>	<i>stx2</i> variants	<i>eae</i> γ	<i>ehxA</i>	PT	ST	Resistance	Disease
154-00	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	2	11		ND
851-00	O157:H7	—	+	<i>stx2</i>	+	+	14	11		BD
1527-00	O157:H7	—	—	<i>stx2</i>	+	+	4	11		HUS
2965-00	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	RDNC	11		BD
1688-01	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	50	11		BD
2110-01	O157:H7	—	—	<i>stx2c</i>	+	+	8	11		BD
1787-02	O157:H7	—	+	<i>stx2</i>	+	+	32	11		HUS
1117-03	O157:H7	—	+	<i>stx2</i>	+	+	32	11		BD, HUS
1357-03	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	32	11		HUS
1899-03	O157:H7	—	+	<i>stx2</i>	+	+	RDNC	11	Te	BD
002-04	O157:[H7]	+	—	<i>stx2</i>	+	+	23	11		BD
256-04	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	49	11		BD
477-04	O157:H7	—	+	<i>stx2</i>	+	+	32	11		ND
774-04	O157:[H7]	+	—	<i>stx2</i>	+	+	23	11		D, HUS
947-04	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	49	11		BD
995-04	O157:[H7]	—	—	<i>stx2</i>	+	+	8	11		BD
1024-04	O157:[H7]	—	—	<i>stx2</i>	+	+	8	11		BD, HUS
1381-04	O157:[H7]	—	+	<i>stx2c</i>	+	+	8	11		BD
1097-05	O157:H7	—	+	<i>stx2</i>	+	+	32	11		BD
1127-05	O157:H7	—	+	<i>stx2</i>	+	+	32	11		BD
1144-05	O157:H7	—	+	<i>stx2</i>	+	+	32	11		ND
1193-05	O157:H7	—	—	<i>stx2</i>	+	+	32	11		ND
1431-05	O157:H7	—	+	<i>stx2</i>	+	+	RDNC	11		ND
1973-05	O157:[H7]	+	—	<i>stx2</i>	+	+	RDNC	11		BD
2188-05	O157:[H7]	—	+	<i>stx2c</i>	+	+	8	11		BD
680-06	O157:H7	—	—	<i>stx2</i>	+	+	8	11		A, D, HUS
746-06	O157:[H7]	—	+	<i>stx2c</i>	+	+	8	11		BD
881-06	O157:H7	—	+	<i>stx2</i>	+	+	32	11		BD, HUS
1206-06	O157:H7	—	+	<i>stx2</i>	+	+	32	11		BD
1346-06	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	71	11	Te	BD
445-07	O157:H7	—	+	<i>stx2</i>	+	+	32	11		ND
645-07	O157:H7	—	+	<i>stx2</i>	+	+	32	11		BD
1478-07	O157:H7	—	—	<i>stx2c</i>	+	+	32	11		BD
1479-07	O157:H7	—	+	<i>stx2</i>	+	+	32	11		A, BD, HUS
1971-07	O157:H7	—	—	<i>stx2</i>	+	+	32	11		A, D, HUS
2298-07	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	49	11		BD
787-08	O157:H7	—	+	<i>stx2</i>	+	+	RDNC	11		BD, HUS
977-08	O157:[H7]	—	+	<i>stx2c</i>	+	+	32	11		BD
1620-08	O157:H7	—	+	<i>stx2</i>	+	+	32	11		A, D, HUS
2248-08	O157:[H7]	—	+	<i>stx2c</i>	+	+	8	11		BD
2437-08	O157:H7	—	+	<i>stx2</i>	+	+	32	11		ND
049-09	O157:H7	—	+	<i>stx2</i>	+	+	32	11		ND
051-09	O157:[H7]	+	—	<i>stx2</i>	+	+	RDNC	11		ND
1402-09	O157:H7	—	—	<i>stx2</i> , <i>stx2c</i>	+	+	49	11		D, HUS

[H7], Negative by latex agglutination serotyping, positive by PCR for *fliC*; PT, phage type; RDNC, Reacts but Does Not Confirm to a published typing pattern; ST, sequence type; Te, tetracycline; A, anaemia; BD, bloody diarrhoea; D, non-bloody diarrhoea; HUS, haemolytic uraemic syndrome.

Antimicrobial susceptibility testing

All O157 STEC strains were susceptible to 11 or more antibiotic agents (AM, AMC, CAZ, CF, CIP, CPD, CTX, CXM, FEP, FOX, GM). Of the 44 strains, two (4.5%) were resistant to Te.

DISCUSSION

The 44 strains investigated in this study are highly representative for Switzerland. This is the total number of STEC O157 pathogens isolated by NENT from all human material submitted over the 10-year period

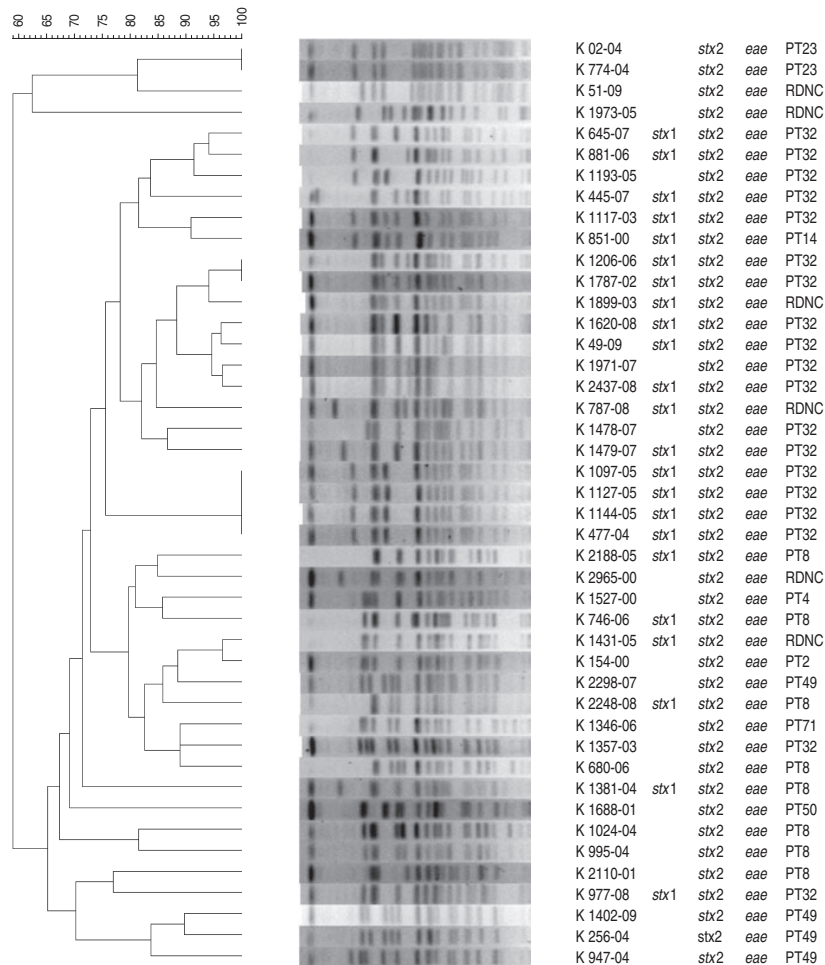


Fig. 2. PFGE dendrogram (Dice similarity index and UPGMA method) of 44 STEC O157 strains isolated from human patients from 2000 to 2009 in Switzerland. PT, Phage type; RDNC, Reacts but Does Not Confirm to a published typing pattern; *stx2*, *stx2* variants (*stx2* and/or *stx2c*).

(2000–2009). Switzerland is a small country in the centre of Europe that is multi-cultural and thus suffers epidemiological influences from several neighbouring ethnic backgrounds. Moreover, the Swiss are known to be the most travelled people in the world. With a population of only 7.1 million, Swiss people execute about 2.2 million trips abroad per year. These aspects also warrant international relevance for our data.

HUS, as a consequence of STEC O157 infection, was recorded in 13 (29.5%) of the 44 investigated patients. Of patients with HUS 84.6% were aged ≤ 5 years. Of the HUS patients, six (46.2%) were male, seven (53.8%) were female. A similar age and gender distribution was reported in a study spanning a 4-year period in Germany [17].

In contrast, in a study performed over a 7-year period (2000–2006) involving 3464 STEC O157 isolates in Northern America, the authors observed fewer HUS cases (6.3%) and fewer HUS patients aged < 5 years (15.3%) [18]. Similar data to these were obtained in Minnesota, Wales and England [1, 19, 20].

Of the 44 STEC O157 strains, 47.7% harboured *stx2* variants only and 52.3% possessed a combination of *stx1* and *stx2* variant genes. No strain was positive for *stx1* only. A similar distribution of *stx* genes was found for clinical STEC O157 isolates in different countries [1, 3, 19–22]. In our study, in all HUS cases strains harbouring the *stx2* gene type were involved. Nevertheless, there was no evidence that the *stx2* variants or the combination of *stx1*, *stx2* and

stx2c genes are directly associated with the clinical outcome.

All 44 STEC O157 strains were positive for *ehxA* and *eae* irrespective of the clinical symptoms. Orth & Wurzner [21] gained comparable results with all isolates investigated in their study being positive for *stx2*, *eae* and *ehxA*. Similar to other studies [3, 23], intimin gamma was the *eae* subtype in our O157:H7 strains.

For years STEC O157 has been classically mentioned as being nSF and motile. In 1990 an outbreak of HUS in Germany, for the first time yielded SF O157:H⁻ [24]. This loss of motility in SF STEC O157 is caused by a 12-bp in-frame deletion in *fliC* that is required for transcriptional activation of genes involved in flagellum biosynthesis [25]. Although a low prevalence of SF O157 strains were found in our strain collection, their occurrence warrants the use of additional methods other than SMAC.

In Germany SF STEC O157:H⁻ are the second most common cause of sporadic HUS [26] and have, up to 2005, caused two large outbreaks involving 57 HUS cases of which seven children died [26].

Among our isolates four (9.1%) SF strains and seven (15.9%) nSF strains were non-motile. HUS developed from 11 classical nSF O157:H7, one atypical SF O157:H⁻ and one nSF O157:H⁻. Of the other three patients with SF STEC O157 isolates, two developed bloody diarrhoea, and from one, no data was available. Patients with SF STEC O157 were aged 1, 2, 5 and 7 years. In Germany, SF STEC O157 was present in 13.3–40.5% of HUS patients, in 7.4–25% of patients with diarrhoea and only in children aged <3 years [27]. All SF STEC O157:H⁻ strains harboured the *fliC* gene-encoding H7 antigen, as in our study.

To further characterize our strains we performed phage-typing and PFGE. Among our STEC O157 isolates nine different phage types were found, with PT8, PT32 and PT49 the most prevalent. In a study on 415 STEC O157 patients in Wales, the most common among 19 detected phage types were PT2 and PT49 showing frequencies of 42.3% and 12.7%, respectively [19]. Similar results were found in an English study [28], where PT2 (46%) was the most common phage type, followed by PT49 (17%) and PT8 (8%). In a recent study from Spain, PT2 was also the most frequently found phage type in human STEC O157, followed by PT8 [22]. Therefore, it appears that Switzerland has a different situation.

PFGE patterns were very heterogeneous except for eight strains (18.2%). Of these eight strains there were two sets each consisting of two strains with similar

patterns (set 1: strain K 02-04, male, aged 1 year, and strain K 774-04, male aged 5 years, both strains isolated in 2004, both *stx2*, *eae* positive and PT23; set 2: strain K 1787-02, male aged 2 years, sample isolated in 2002, and strain K 1206-06, female aged 45 years, sample isolated in 2006, both *stx1*, *stx2*, *eae* positive and PT32) and one set consisting of four equal patterns, all four PT32 and *stx1*, *stx2* and *eae* positive (K 447-04, female aged 10 years, sample isolated in 2004 and K 1097-05, male aged 2 years, K 1127-05, male aged 7 years and K 1144-05, female aged 40 years, all three samples isolated in 2005). The high genetic diversity within the strains leads to the conclusion that STEC O157 infections in Switzerland most often occur as single cases. Similar to the situation in Switzerland, only a few STEC O157:H7-associated outbreaks have been reported in the neighbouring countries within this time period [29–31].

In our study a very low prevalence of antibiotic resistance was found for STEC O157 isolates. Only two strains (4.5%) were resistant to tetracycline. In England and Wales 20% of human STEC O157 strains collected between 1995 and 1998 were resistant to one or more antimicrobial agents [20]. Only 1% of human STEC showed resistance in Germany [32]. Recently, Srinivasan *et al.* [33] investigated 153 STEC O157:H7/H⁻ strains isolated from human faeces, cows and food in the USA of which >90% showed resistance to ampicillin and cephalothin. Moreover, resistance to tetracycline, gentamicin, cefotaxime and ciprofloxacin was found. High prevalence of resistance was also found in O157:H7 strains from Spanish people [22], where 38% and 24% of the isolates showed resistance to tetracycline and ampicillin, respectively.

To summarize, only 30.6% of the STEC strains isolated from clinical cases from 2000 to 2009 in Switzerland were STEC O157. SF and nSF strains were found. All strains were positive for *stx2* variants (*stx2* and/or *stx2c*), *eae* and *ehxA* and showed a very favourable antibiotic resistance situation. The high genetic diversity within the strains leads to the conclusion that STEC O157 infections in Switzerland most often occur as sporadic cases.

ACKNOWLEDGEMENTS

We thank the FOPH (Federal Office of Public Health) for support in collecting the medical history data and Grethe Sägerser for help in strain collection and technical support.

DECLARATION OF INTEREST

None.

REFERENCES

1. Hedican EB, *et al.* Characteristics of O157 versus non-O157 shiga toxin-producing *Escherichia coli* infections in Minnesota, 2000–2006. *Clinical Infectious Disease* 2009; **49**: 358–364.
2. Werber D, *et al.* Shiga toxin-producing *Escherichia coli* serogroups in food and patients, Germany. *Emerging Infectious Disease* 2008; **14**: 1803–1806.
3. Blanco JE, *et al.* Serotypes, virulence genes, and intimin types of shiga toxin (verotoxin)-producing *Escherichia coli* isolates from human patients: prevalence in Lugo, Spain, from 1992 through 1999. *Journal of Clinical Microbiology* 2004; **42**: 311–319.
4. Feng PC, *et al.* Genetic diversity among clonal lineages within *Escherichia coli* O157:H7 stepwise evolutionary model. *Emerging Infectious Disease* 2007; **13**: 1701–1706.
5. Bielaszewska M, *et al.* Isolation and characterization of sorbitol-fermenting shiga toxin (verocytotoxin)-producing *Escherichia coli* O157:H[−] strains in the Czech Republic. *Journal of Clinical Microbiology* 1998; **36**: 2135–2137.
6. Friedrich AW, *et al.* Prevalence, virulence profiles, and clinical significance of shiga toxin-negative variants of enterohemorrhagic *Escherichia coli* O157 infection in humans. *Clinical Infectious Disease* 2007; **45**: 39–45.
7. Bielaszewska M, *et al.* Shiga toxin-negative attaching and effacing *Escherichia coli*: distinct clinical associations with bacterial phylogeny and virulence traits and inferred in-host pathogen evolution. *Clinical Infectious Disease* 2008; **47**: 208–217.
8. Zweifel C, Stephan R. Microbiological monitoring of sheep carcass contamination in three Swiss abattoirs. *Journal of Food Protection* 2003; **66**: 946–952.
9. Abdulmawjood A, Roth S, Bulte M. Two methods for construction of internal amplification controls for the detection of *Escherichia coli* O157 by polymerase chain reaction. *Molecular and Cellular Probes* 2002; **16**: 335–339.
10. Gannon VP, *et al.* Use of the flagellar H7 gene as a target in multiplex PCR assays and improved specificity in identification of enterohemorrhagic *Escherichia coli* strains. *Journal of Clinical Microbiology* 1997; **35**: 656–662.
11. Russmann H, *et al.* Genotyping of shiga-like toxin genes in non-O157 *Escherichia coli* strains associated with haemolytic uraemic syndrome. *Journal of Medical Microbiology* 1995; **42**: 404–410.
12. Blanco M, *et al.* Serotypes, virulence genes, and intimin types of shiga toxin (verotoxin)-producing *Escherichia coli* isolates from healthy sheep in Spain. *Journal of Clinical Microbiology* 2003; **41**: 1351–1356.
13. Schmidt H, Beutin L, Karch H. Molecular analysis of the plasmid-encoded hemolysin of *Escherichia coli* O157:H7 strain EDL 933. *Infection and Immunity* 1995; **63**: 1055–1061.
14. Stephan R, Hoelzle LE. Characterization of shiga toxin type 2 variant B-subunit in *E. coli* strains from asymptomatic human carriers by PCR-RFLP. *Letters in Applied Microbiology* 2000; **31**: 139–142.
15. Khakhria R, Duck D, Lior H. Extended phage-typing scheme for *Escherichia coli* O157:H7. *Epidemiology and Infection* 1990; **105**: 511–520.
16. Wirth T, *et al.* Sex and virulence in *Escherichia coli*: an evolutionary perspective. *Molecular Microbiology* 2006; **60**: 1136–1151.
17. Ammon A, Petersen LR, Karch H. A large outbreak of hemolytic uremic syndrome caused by an unusual sorbitol-fermenting strain of *Escherichia coli* O157:H[−]. *Journal of Infectious Diseases* 1999; **179**: 1274–1277.
18. Gould LH, *et al.* Hemolytic uremic syndrome and death in persons with *Escherichia coli* O157:H7 infection, foodborne diseases active surveillance network sites, 2000–2006. *Clinical Infectious Disease* 2009; **49**: 1480–1485.
19. Chalmers RM, *et al.* The surveillance of vero cytotoxin-producing *Escherichia coli* O157 in Wales, 1990 to 1998. *Emerging Infectious Disease* 1999; **5**: 566–569.
20. Willshaw GA, *et al.* Verocytotoxin-producing *Escherichia coli* (VTEC) O157 and other VTEC from human infections in England and Wales: 1995–1998. *Journal of Medical Microbiology* 2001; **50**: 135–142.
21. Orth D, Wurzner R. What makes an enterohemorrhagic *Escherichia coli*? *Clinical Infectious Disease* 2006; **43**: 1168–1169.
22. Mora A, *et al.* Antimicrobial resistance of shiga toxin (verotoxin)-producing *Escherichia coli* O157:H7 and non-O157 strains isolated from humans, cattle, sheep and food in Spain. *Research in Microbiology* 2005; **156**: 793–806.
23. Beutin L, *et al.* Characterization of shiga toxin-producing *Escherichia coli* strains isolated from human patients in Germany over a 3-year period. *Journal of Clinical Microbiology* 2004; **42**: 1099–1108.
24. Karch H, *et al.* Verocytotoxin-producing *Escherichia coli* causing a hemolytic-uremic syndrome in children. *Deutsche Medizinische Wochenschrift* 1990; **115**: 489–495.
25. Monday SR, Minnich SA, Feng PC. A 12-base-pair deletion in the flagellar master control gene *fliC* causes nonmotility of the pathogenic German sorbitol-fermenting *Escherichia coli* O157:H[−] strains. *Journal of Bacteriology* 2004; **186**: 2319–2327.
26. Karch H, Tarr PI, Bielaszewska M. Enterohaemorrhagic *Escherichia coli* in human medicine. *International Journal of Medical Microbiology* 2005; **295**: 405–418.
27. Karch H, Bielaszewska M. Sorbitol-fermenting shiga toxin-producing *Escherichia coli* O157:H(−) strains: epidemiology, phenotypic and molecular characteristics, and microbiological diagnosis. *Journal of Clinical Microbiology* 2001; **39**: 2043–2049.
28. Thomas A, *et al.* Vero cytotoxin-producing *Escherichia coli*, particularly serogroup O157, associated with

- human infections in England and Wales: 1992–1994. *Epidemiology and Infection* 1996; **117**: 1–10.
29. **Alpers K, et al.** Sorbitol-fermenting enterohaemorrhagic *Escherichia coli* O157:H[−] causes another outbreak of haemolytic uraemic syndrome in children. *Epidemiology and Infection* 2009; **137**: 389–395.
30. **Conedera G, et al.** A family outbreak of *Escherichia coli* O157 haemorrhagic colitis caused by pork meat salami. *Epidemiology and Infection* 2007; **135**: 311–314.
31. **King L, et al.** Community-wide outbreak of *Escherichia coli* O157:H7 associated with consumption of frozen beef burgers. *Epidemiology and Infection* 2009; **137**: 889–896.
32. **Schmidt H, et al.** Antibiotic susceptibilities of verocytotoxin-producing *Escherichia coli* O157 and non-O157 strains isolated from patients and healthy subjects in Germany during 1996. *Journal of Antimicrobial Chemotherapy* 1998; **42**: 548–550.
33. **Srinivasan V, et al.** Antimicrobial resistance patterns of shiga toxin-producing *Escherichia coli* O157:H7 and O157:H7[−] from different origins. *Microbial Drug Resistance* 2007; **13**: 44–51.

3. Danksagung

An dieser Stelle möchte ich allen danken, die zum Gelingen dieser Arbeit beigetragen haben.

Besonderer Dank geht an:

Herrn Prof. Dr. Roger Stephan, Institut für Lebensmittelsicherheit und -hygiene, Vetsuisse Fakultät Universität Zürich, für die Möglichkeit zur Erstellung dieser Dissertation, die fachliche Beratung, das grosszügige Entgegenkommen in allen Belangen, sowie die Übernahme des Referates. Ganz besonders möchte ich ihm danken für seine offene, liebe Art. Stets konnte ich auf ihn zählen und er stand mir mit Rat und Tat zur Seite.

Herrn PD Dr. Lothar Beutin, Nationales Referenzlabor für Escherichia coli (NRL-E.coli), Bundesinstitut für Risikobewertung (BfR) Berlin, für die Übernahme des Korreferates.

Herrn PD Dr. Herbert Hächler, Institut für Lebensmittelsicherheit und -hygiene, Nationales Zentrum für enteropathogene Bakterien (NENT), für die Zusammenarbeit bei dieser Dissertation, die fachliche Betreuung, die vielen interessanten und lehrreichen Gespräche und Erklärungen. Besonders aber möchte ich ihm danken für die tolle Zeit als Zimmernachbarn, die vielen gemeinsamen Lacher, dass er sich immer Zeit für mich genommen hat, mir alles ausführlich erklärt hat, mich stets ermuntert hat und mir zur Seite stand.

Das BAG, mit Hilfe dessen wir die anamnestischen Daten ergänzen konnten.

Frau Grethe Sägesser, für die Hilfe beim Sammeln der Stämme und ihre frohe, belebende Natur.

Frau Nicole Cernela-Giezendanner, für die kompetente Einführung in die Laborarbeit, die grosse Mithilfe bei der Charakterisierung unserer Stämme und Rat und Tat in jedem Belangen. Für ihre offene, liebe Art und die Freundschaft, die sie mir entgegen gebracht hat, möchte ich mich aber am meisten bedanken. Ich bin froh, dass wir uns kennen gelernt haben.

Meine Mitdissertantinnen, für unzählige freundschaftliche und aufheiternde Gespräche.

Das gesamte ILS-Team für die angenehme Atmosphäre am Institut, die Materialien Vor-, Zu-, und Nachbereitung, die netten und interessanten Gespräche und gute Zusammenarbeit.

Meine Freunde für die offenen Arme und Ohren während meiner ganzen Ausbildungszeit und dass sie mir immer wieder Energie und Mut gegeben haben zum Weitermachen.

Meine Eltern, die mir diesen tollen Weg ermöglicht und mich dabei begleitet haben. Ohne Euch wäre das alles nie möglich gewesen. Vielen herzlichen Dank.

Meine Schwester, die sich meine Sorgen und Geschichten angehört hat, mir zur Seite stand und Ihrer bezaubernden Familie, die mich aus jedem Tief holen konnte. Hab Dich fest lieb.

Meine grosse Liebe. Diego, ohne Dich wäre gar nichts möglich, Du gibst mir Kraft und Energie für alles, was ich angehe. Du hast mich stets unterstützt und mir geholfen meinen Weg zu gehen. Durch Deine Liebe bin ich. Ich liebe Dich über alles. Danke.

4. Curriculum Vitae

Name, Vornamen	Käppeli, Ursula Elisabeth
Geburtsdatum	22.09.1983
Geburtsort	Muri, Aargau
Nationalität	CH
Heimatort	Wohlen, Aargau
08/1990 - 07/1995	Primarschule, Wohlen, Aargau, Schweiz
08/1995 - 07/1999	Bezirksschule, Wohlen, Aargau, Schweiz
08/1999 - 05/2003	Kantonsschule, Wohlen, Aargau, Schweiz
28.06.2003	Matura, Kantonsschule, Wohlen, Aargau, Schweiz
09/2004 - 10/2009	Studium der Veterinärmedizin, Universität Zürich, Zürich, Schweiz
15.10.2009	Abschlussprüfung vet. med., Universität Zürich, Zürich, Schweiz
11/2009 - 07/2010	Anfertigung der Dissertation unter Leitung von Prof. Dr. Roger Stephan am Institut für Lebensmittelsicherheit und -hygiene der Vetsuisse-Fakultät, Universität Zürich
11/2009 - 08/2010	Doktorandin, Institut für Lebensmittelsicherheit und -hygiene, Vetsuisse-Fakultät, Universität Zürich, Zürich, Schweiz